DNA Methylation Analysis (Agilent's Genomic Workbench Steps)

Standard Procedure for Analysis

Project Name: DNA methylation Analysis

Requestor:

Names and contact info of Analyst:

Brief Description of Analysis Types: DNA methylation tilling array data

Procedure or Analysis Revision Date:

Analysis Results/Deliverable Date:

Scientific context or hypothesis for analysis:

To access differential methylation status among different conditions

Checklist before starting analysis:

Scanned DNA tilling array chips Experiment design Sample annotation Sources of input required:

- Experiment design (Control, Disease, Sampling time point),
- Sample annotation,
- Tissue and other supporting information

Input data:

Agilent Two Color tilling array feature extracted txt files

Analysis plan or steps taken:

1. Purpose.

To analyze DNA methylation microarray data generated using Agilent's ChIP-on-Chip tilling array platform

2. Responsibility.

Systems Biology

3. Materials and Equipment.

Genomic Workbench v 7.0.4 Lit Edition Software (Agilent Technology Inc.)

4. Procedure.

Recently, we use R for dealing with Agilent DNA methylation microarrays. The preprocessing steps are same to the transcriptomics analysis.

The downstream analysis is typically divided into specific regions.

Data import and Experiment Set up

To set up an experiment:

- Select the CH3 application
- Import design files and data files (Agilent FE files; Import > Array Files > FE File)
- Create a new experiment
- · Add the imported data to the experiment
- Select the experiment
- Combine design and generate report

To update probe annotation in design files

- Home tab, >User Preferences > Miscellaneous tab > eArray User Details (type Username and Password) > OK
- select the appropriate design > right- click the desired design > Download from eArray.com > Yes

To generate a probe report

- Reports tab > Generate Report > Complete Genome > type the report name (name.xls) > OK.
- The probe report includes the following information either for single chromosomes or the complete genome:
- Log ratio for each event
- Number of probes
- Start and stop of each event
- Whether the probe is in a CpG island, a promoter, or inside a gene boundary
- Probable gene for the probe
- Excel spreadsheet matrix containing all the data (and the above probes and samples annotations).

5. References.

Genomic Workbench 7.0.4 Methylation (CH3) Analysis User Manual (G3800-90010_MethylationAnalysis.pdf)
http://www.r-project.org/